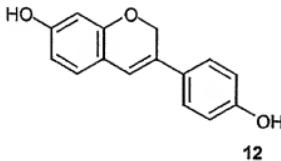


AMENDMENTS TO THE CLAIMS

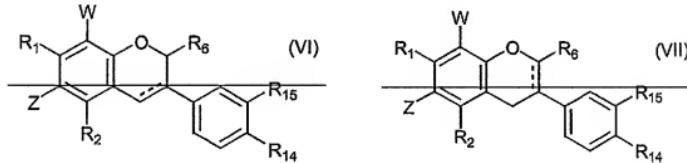
This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (currently amended): A method of increasing the sensitivity of cancer cells or a tumour to a chemotherapeutic agent by contacting said cells or tumour with an isoflavanoid compound of formula 12(VI) or (VII):



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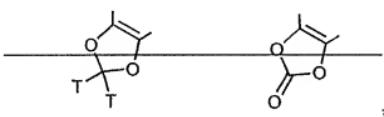
wherein

R₁, R₂ and Z are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₂ is as previously defined, and R₁ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



, or

R_4 is as previously defined, and R_2 and Z taken together with the carbon atoms to which they are attached form a five membered ring selected from



,

W is R_1 ,

R_3 is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, $C(O)R_{11}$ where R_{11} is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO_2R_{12} where R_{12} is hydrogen, alkyl, haloalkyl, aryl or arylalkyl,

R_4 is hydrogen, alkyl or aryl, or

R_3 and R_4 taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl;

R_6 is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR_3R_4 , COR_{11} where R_{11} is as previously defined, CO_2R_{12} where R_{12} is as previously defined or $CONR_3R_4$,

R_9 is alkyl, haloalkyl, aryl, arylalkyl, $C(O)R_{11}$ where R_{11} is as previously defined, or $Si(R_{13})_3$ where R_{13} where each R_{13} is independently hydrogen, alkyl or aryl,

R_{10} is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamine;

the drawing “—” represents either a single bond or a double bond;

T is independently hydrogen, alkyl or aryl,

R₁₄, and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀,

COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure;

and pharmaceutically acceptable salts thereof, and,

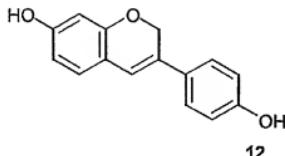
wherein the cancer is ovarian, pancreatic or prostate cancer, and

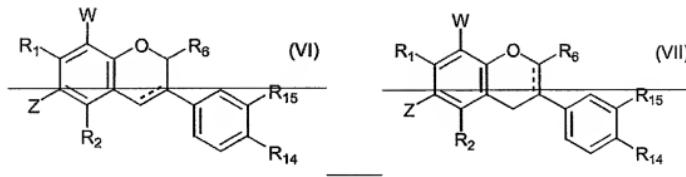
the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicin.

2. (previously presented): A method of claim 1, wherein prior to the contacting, the cancer cells or tumour were/was not sensitive to the chemotherapeutic agent.

3. (currently amended): A method of claim 1, wherein the compound of formula 12(VI)-or-(VII) is administered to a subject in need of such treatment.

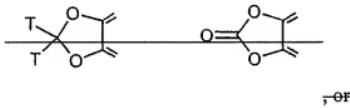
4. (currently amended): A combination therapy for the treatment or prophylaxis of cell proliferation, cancer or a disease associated with oxidant stress comprising administering to a subject a therapeutically effective amount of a compound of formula 12(VI)-or-(VII) and a chemotherapeutic agent:



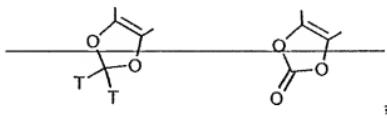


wherein

R_1 , R_2 and Z are independently hydrogen, hydroxy, OR_9 , $OC(O)R_{10}$, $OS(O)R_{10}$, CHO , $C(O)R_{10}$, $COOH$, CO_2R_{10} , $CONR_3R_4$, alkyl, haloalkyl, arylalkyl, arylalkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R_3 is as previously defined, and R_1 and Z taken together with the carbon atoms to which they are attached form a five membered ring selected from



R_1 is as previously defined, and R_2 and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



W is R₁,

R₃ is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R₁₁ where R₁₁ is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO₂R₁₂ where R₁₂ is hydrogen, alkyl, haloalkyl, aryl or arylalkyl,

R₄ is hydrogen, alkyl or aryl, or

R₃ and R₄ taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl,

R₆ is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR₃R₄, COR₁₁ where R₁₁ is as previously defined, CO₂R₁₂ where R₁₂ is as previously defined or CONR₃R₄,

R₉ is alkyl, haloalkyl, aryl, arylalkyl, C(O)R₁₁ where R₁₁ is as previously defined, or Si(R₁₃)₃ where R₁₃ where each R₁₃ is independently hydrogen, alkyl or aryl,

R₁₀ is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino,

the drawing “—” represents either a single bond or a double bond,

T is independently hydrogen, alkyl or aryl,

R₁₄, and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure,

and pharmaceutically acceptable salts thereof, and,

wherein the cancer is ovarian, pancreatic or prostate cancer, and

the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicin.

5.-7. (canceled).

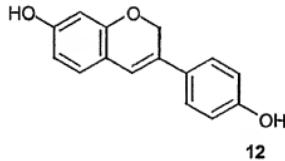
8. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) or (VII) precedes the administration of the chemotherapeutic agent.

9. (currently amended): A method of claim 4, wherein the administration of the compound of formula 12(VI) or (VII) and the chemotherapeutic agent is simultaneous.

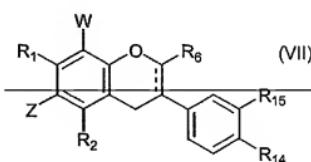
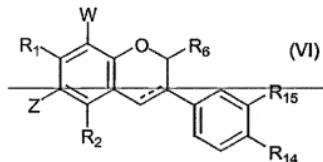
10. (currently amended): A method claim 4, wherein the combination therapy follows observed resistance by cancer cells or tumour to [[a]]the chemotherapeutic agent.

11,-22, (canceled),

23. (currently amended): A pharmaceutical composition comprising a compound of formula 12(VI) or (VII) and a chemotherapeutic agent:

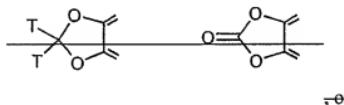


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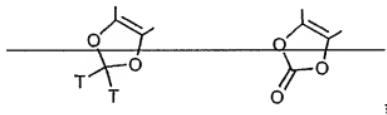
wherein

R₁, R₂ and Z are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, alkylaryl, alkoxyaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₂ is as previously defined, and R₁ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



, or

R₁ is as previously defined, and R₂ and Z taken together with the carbon atoms to which they are attached form a five-membered ring selected from



,

W is R₄

R₃ is hydrogen, alkyl, arylalkyl, alkenyl, aryl, an amino acid, C(O)R₁₁ where R₁₁ is hydrogen, alkyl, aryl, arylalkyl or an amino acid, or CO₂R₁₂ where R₁₂ is hydrogen, alkyl, haloalkyl, aryl or arylalkyl;

R₄ is hydrogen, alkyl or aryl, or

R₃ and R₄ taken together with the nitrogen to which they are attached comprise pyrrolidinyl or piperidinyl;

R₆ is hydrogen, hydroxy, alkyl, aryl, amino, thio, NR₃R₄, COR₁₁ where R₁₁ is as previously defined, CO₂R₁₂ where R₁₂ is as previously defined or CONR₃R₄,

~~R₉ is alkyl, haloalkyl, aryl, arylalkyl, C(O)R₁₁ where R₁₁ is as previously defined, or Si(R₁₃)₃, where R₁₃ where each R₁₃ is independently hydrogen, alkyl or aryl,~~

~~R₁₀ is hydrogen, alkyl, haloalkyl, amino, aryl, arylalkyl, an amino acid, alkylamino or dialkylamino;~~

~~the drawing “—” represents either a single bond or a double bond;~~

~~T is independently hydrogen, alkyl or aryl,~~

~~R₁₄, and R₁₅ are independently hydrogen, hydroxy, OR₉, OC(O)R₁₀, OS(O)R₁₀, CHO, C(O)R₁₀, COOH, CO₂R₁₀, CONR₃R₄, alkyl, haloalkyl, arylalkyl, alkenyl, alkynyl, aryl, heteroaryl, thio, alkylthio, amino, alkylamino, dialkylamino, nitro or halo, or R₁₄ and R₁₅ are fused together to form a cyclic alkyl, aromatic or heteroaromatic structure,~~

~~and pharmaceutically acceptable salts thereof, and,~~

~~wherein the chemotherapeutic agent is platinum-based or anti-mitotic agent cisplatin, carboplatin, paclitaxel, gemcitabine or doxorubicins.~~

24.-28. (Canceled).